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Applicants Wilson et. al.	Examiner Murphy
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U.S. PATENT DOCUMENTS

*Exam. Init.	No.	Document No.							Date	Name	Class	Subclass	Filing Date if Appropriate

FOREIGN PATENT DOCUMENT

		Document No.									Class	SubClass	Translator Yes No

OTHER DOCUMENTS (including Author, Title Date, Pertinent Pages, Etc.)

A	Peral et al., 1995, "Splicing mutations of the polycystic kidney disease 1 (PKD1) gene induced by intronic deletion," <i>Human Molec. Genet.</i> 4:569-574.
B	The European Polycystic Kidney Disease Consortium, 1994, "The polycystic kidney disease 1 gene encodes a 14kb transcript and lies within a duplicated region on chromosome 16," <i>Cell</i> 77:881-894.
C	Watnick et al., 1997, "An unusual pattern of mutation in the duplicated portion of PKD1 is revealed by use of a novel strategy for mutation detection," <i>Human Molecular Genetics</i> 6:1473-1481.
D	Longa et al., 1997, "A large TSC2 and PKD1 gene deletion is associated with renal and extrarenal signs of autosomal dominant polycystic kidney disease," <i>Nephrology Dialysis Transplantation</i> 12:1900-1907
E	Thomas et al., 1999, "Identification of mutations in the repeated part of the autosomal dominant polycystic kidney disease type 1 gene, PKD1, by long-range PCR," <i>Am. J. Human Genetics</i> 65:39-49.